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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
10/825,457	04/14/2004	Chih-Ping Liu	55600-8014.US02	8343
22918 PERKINS COI	7590 03/26/200 E LLP	EXAMINER		
P.O. BOX 2168		DANG, IAN D		
MENLO PARK	x, CA 94026		ART UNIT	PAPER NUMBER
			1647	
			MAIL DATE	DELIVERY MODE
			03/26/2008	PAPER

Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

Advisory Action Before the Filing of an Appeal Brief

Application No.	Applicant(s)	
10/825,457	LIU ET AL.	
Examiner	Art Unit	
IAN DANG	1647	

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The MAILING DATE of this communication appe	ars on the cover sheet with the d	correspondence add	ress
THE REPLY FILED <u>19 February 2008</u> FAILS TO PLACE THIS A	APPLICATION IN CONDITION FO	R ALLOWANCE.	
 The reply was filed after a final rejection, but prior to or on application, applicant must timely file one of the following application in condition for allowance; (2) a Notice of Apperfor Continued Examination (RCE) in compliance with 37 C periods: 	the same day as filing a Notice of A replies: (1) an amendment, affidavited al (with appeal fee) in compliance	Appeal. To avoid abar t, or other evidence, w with 37 CFR 41.31; or	hich places the (3) a Request
a) The period for reply expires 4 months from the mailing date b) The period for reply expires on: (1) the mailing date of this Ar no event, however, will the statutory period for reply expire to Examiner Note: If so checked, check either box (a) or (4)	dvisory Action, or (2) the date set forth in ater than SIX MONTHS from the mailing b). ONLY CHECK BOX (b) WHEN THE	g date of the final rejection	n.
MONTHS OF THE FINAL REJECTION. See MPEP 706.07(f Extensions of time may be obtained under 37 CFR 1.136(a). The date of have been filed is the date for purposes of determining the period of extunder 37 CFR 1.17(a) is calculated from: (1) the expiration date of the set forth in (b) above, if checked. Any reply received by the Office later may reduce any earned patent term adjustment. See 37 CFR 1.704(b). NOTICE OF APPEAL	on which the petition under 37 CFR 1.1 ension and the corresponding amount of hortened statutory period for reply origi	of the fee. The appropria nally set in the final Offic	ate extension fee e action; or (2) as
 The Notice of Appeal was filed on A brief in comp filing the Notice of Appeal (37 CFR 41.37(a)), or any exter Notice of Appeal has been filed, any reply must be filed with the properties. 	nsion thereof (37 CFR 41.37(e)), to	avoid dismissal of the	
AMENDMENTS			
3. The proposed amendment(s) filed after a final rejection, by (a) They raise new issues that would require further core.	nsideration and/or search (see NOT		cause
 (b) ☐ They raise the issue of new matter (see NOTE below (c) ☐ They are not deemed to place the application in better appeal; and/or 	•	ducing or simplifying th	ne issues for
(d) ☐ They present additional claims without canceling a c	corresponding number of finally reje	ected claims.	
NOTE: (See 37 CFR 1.116 and 41.33(a)).	21 Con attached Nation of Nan Con		OTOL 224\
 The amendments are not in compliance with 37 CFR 1.12 Applicant's reply has overcome the following rejection(s): 	·		
 Newly proposed or amended claim(s) would be all non-allowable claim(s). 	·	•	_
7. For purposes of appeal, the proposed amendment(s): a) [how the new or amended claims would be rejected is prov The status of the claim(s) is (or will be) as follows:		l be entered and an ex	৻planation of
Claim(s) allowed: Claim(s) objected to:			
Claim(s) rejected: <u>1.3 and 4</u> . Claim(s) withdrawn from consideration: AFFIDAVIT OR OTHER EVIDENCE			
 The affidavit or other evidence filed after a final action, but because applicant failed to provide a showing of good and was not earlier presented. See 37 CFR 1.116(e). 			
9. The affidavit or other evidence filed after the date of filing a entered because the affidavit or other evidence failed to or showing a good and sufficient reasons why it is necessary	vercome <u>all</u> rejections under appea and was not earlier presented. Se	al and/or appellant fails see 37 CFR 41.33(d)(1)	s to provide a).
10.	n of the status of the claims after er	ntry is below or attache	ed.
 The request for reconsideration has been considered but See Continuation Sheet. 	does NOT place the application in	condition for allowand	ce because:
12. ☐ Note the attached Information <i>Disclosure Statement</i>(s). (13. ☐ Other:	PTO/SB/08) Paper No(s)		
	/Robert Landsman/ Primary Examiner, Art U	nit 1647	
	-		

Continuation of 11. does NOT place the application in condition for allowance because:

Applicants' response and arguments filed on 02/19/2008 have overcome the rejection of claims 1, 3, and 4 under 35 USC 112, First paragraph (new matter). The rejection of claims 1, 3, and 4 under 35 USC 112, First paragraph (new matter) has been withdrawn.

Applicant's response, arguments, and amendment made to claim 1 filed 02/19/2008 have been overcome the rejection of claims 1, 3, and 4 under 35 USC 112, First paragraph (Written Description). The references provided by Applicant in Exhibits 1 and 2 disclose that regions required for antiviral activity are known (see page 137 of Pontzer et al.) and conserved regions for interferon tau (see Radhakrishnan et al., paragraph bridging the columns of page 155, especially right column). The rejections of claims 1, 3, and 4 under 35 USC 112, First paragraph (Written Description) has been withdrawn.

The rejection of claims 1, 3, and 4 under USC 112, First paragraph (Enablement) is maintained. Applicant's response and amendment made to claim 1 filed 02/19/2008 have been considered but are not found persuasive regarding the rejections of claims 1, 3, and 4. Applicants are not enabled for a method for treating a patient with multiple sclerosis by reducing IFN-gamma blood levels in a subject comprising orally administering an IFN-tau having greater than about 90% sequence identity to SEQ ID NO:2 at a dosage of between 6 x108 - 5x1012 units to decrease the subject's IFN-tau blood level relative to the IFN-gamma blood level in the absence of IFN-tau administration because Applicants have not provided any guidance on how IFN-tau acts in patients and whether the regions of IFN-tau responsible for antiviral activity would be required or responsible for decreasing IFN-gamma blood levels for the treatment of multiple sclerosis. While the the references provided by Applicant in Exhibits 1 and 2 disclose that regions of IFN-tau required for antiviral activity (see page 137 of Pontzer et al.) and that conserved regions of IFN-tau are known (see Radhakrishnan et al., paragraph bridging the columns of page 155, especially right column), Applicants have not provided any correlation between the antiviral and conserved regions of IFN-tau and their activities in reducing IFN-gamma blood levels in patient for the treatment of MS. If Applicants can provide evidence that the antiviral and conserved regions of IFN-tau disclosed in the references are responsible for the reduction in IFN-gamma blood levels in MS patients, Applicant would be enabled for the claimed method.